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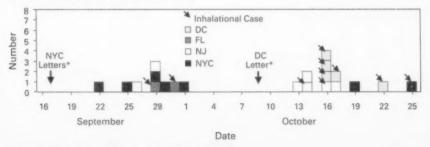
Update: Investigation of Bioterrorism-Related Anthrax and Interim Guidelines for Clinical Evaluation of Persons with Possible Anthrax

Since October 3, 2001, CDC and state and local public health authorities have been investigating cases of bioterrorism-related anthrax. This report updates findings as of October 31, and includes interim guidelines for the clinical evaluation of persons with possible anthrax. A total of 21 cases (16 confirmed and five suspected) of bioterrorism-related anthrax have been reported among persons who worked in the District of Columbia, Florida, New Jersey, and New York City (Figure 1). Until the source of these intentional exposures is eliminated, clinicians and laboratorians should be alert for clinical evidence of *Bacillus anthracis* infection. Epidemiologic investigation of these cases and surveil-lance to detect new cases of bioterrorism-associated anthrax continues.

New York

To date, the investigations in New York City have identified one confirmed inhalational case and six (three confirmed and three suspected) cutaneous anthrax cases; the confirmed inhalational and one suspected cutaneous case have been identified since the last report (1). The six cutaneous cases were associated with four media companies (A–D); the most recent suspected cutaneous case is associated with company D. The most

FIGURE 1. Number of bioterrorism-related anthrax cases, by date of onset and work location — District of Columbia (DC), Florida (FL), New Jersey (NJ), and New York City (NYC), September 16–October 25, 2001



* Postmarked date of known contaminated letters.

recent confirmed inhalational case is not directly associated with any media company or with mail handling. No cases among postal workers have been identified.

The most recent suspected cutaneous case occurred in a 34-year-old man who worked in the mail room of company D who might have handled a letter postmarked September 18, which the patient handled during October 12–15 and subsequently was found to contain *B. anthracis* (1). On October 19, the patient noted a small, erythematous pruritic papule on his left forearm that later developed a small vesicle. On October 21, he started ciprofloxacin. By October 22, an eschar had developed, increased in size, and over the next several days was surrounded by erythema, edema, and induration. A biopsy was positive for *B. anthracis* by immunohistochemical (IHC) staining.

The inhalational anthrax case occurred in a 61-year-old woman who worked in the stockroom of a hospital in Manhattan. The patient became ill on October 25 with malaise and myalgias. During the next several days, she had shortness of breath, chest discomfort, and a productive cough with blood-tinged sputum. She reported no fever, chills, or night sweats. She presented to an emergency department on October 28 in respiratory distress. Her temperature was 102 F (39 C), and she was admitted to the intensive care unit and required mechanical ventilation. Initial chest radiograph revealed pulmonary venous congestion and bilateral pleural effusions; a chest computerized tomography (CT) scan revealed a widened mediastinum and bilateral pleural effusions. An echocardiogram indicated a small pericardial effusion. She was empirically treated with levofloxacin, rifampin, and clindamycin. Blood cultures grew *B. anthracis* less than 24 hours after admission. Her pleural effusion revealed hemorrhagic fluid and *B. anthracis*. The patient died on October 31.

New Jersey

To date, investigations in New Jersey and Pennsylvania have identified seven (five confirmed and two suspected) anthrax cases. Since the last report (1), cutaneous disease was confirmed in two patients, and inhalational anthrax was confirmed in two patients, one of whom was previously classified as a suspected case-patient. Five patients worked in New Jersey at one of two postal facilities. Although no specific contaminated letter was implicated in these cases, contaminated letters destined for both New York City and the District of Columbia passed through at least one of the postal facilities in New Jersey.

Inhalational anthrax was confirmed in a 56-year-old female postal worker who initially was classified as a suspected case-patient (1). Her pleural fluid was positive for *B. anthracis* by polymerase chain reaction (PCR) and a pleural biopsy was positive for *B. anthracis* by IHC staining.

On October 13, a 54-year-old Delaware resident who worked as a mail sorter at a New Jersey postal processing and distributing center developed a painless lesion on the dorsum of his left hand. The lesion began as an erythematous "knot" several millimeters in size that developed a crusted scale during the next few days. No associated edema, eschar, or lymphadenopathy was observed. The patient had elevated levels of serum antibody (lgG) to the protective antigen component of the anthrax toxin using enzyme-linked immunosorbant assay.

On October 15, a 43-year-old female postal worker who worked at a facility in which anthrax cases have been documented developed fever, headache, chills, and shortness of breath. She was treated with levofloxacin, but her symptoms progressed and she was admitted to a hospital on October 18. A chest radiograph indicated a right perihilar

infiltrate and a small pleural effusion. She was started on multidrug therapy, including ciprofloxacin, which was changed to azithromycin after 24 hours. On admission, she was febrile and tachycardic. She had an elevated white blood cell (WBC) count of 11,000 with 14% bands. A CT scan on October 19 showed a right pleural effusion, perihilar consolidation, and mediastinal adenopathy. She subsequently had two thoracenteses that produced serosanguinous pleural fluid and a bronchoscopy that showed grossly edematous bronchi. Both pleural fluid and bronchial biopsy were positive for *B. anthracis* by IHC stain.

On October 17, a 51-year-old woman developed a large pimple on her forehead with erythema and swelling. On October 18, the lesion enlarged, was slightly painful, nonpruritic, and drained a small amount of yellowish fluid. She sought medical care, cervical and preauricular lymphadenopathy was noted on physical examination, and she was treated with ciprofloxacin. The lesion progressed and ulcerated. On October 22, she presented to an emergency department and was admitted with a diagnosis of cellulitis. On admission, she was afebrile with normal vital signs and had a swollen right face and eyelid and enlarged right anterior cervical nodes. Intravenous ciprofloxacin for cutaneous anthrax was started. On October 24, the ulcer was biopsied and debrided. Biopsy specimens were positive for *B. anthracis* by PCR and IHC. The patient improved and was discharged on October 27 on oral ciprofloxacin. The patient worked as a bookkeeper and reported receiving no unusual or powder-containing mail at home or work. She had made no visits to any post offices in several months.

District of Columbia

To date, investigations in the District of Columbia, Maryland, and Virginia have confirmed inhalational anthrax in four persons who worked at one postal facility in the District of Columbia. An additional case of inhalational anthrax has been confirmed in a 59-year-old postal worker in a U.S. State Department mail sorting facility that receives mail from the District of Columbia postal facility associated with the previous four cases. The patient presented to an emergency department on October 24 with temperature of 100.8 F (38 C), sweats, myalgia, chest discomfort, mild cough, nausea, vomiting, diarrhea, and abdominal pain. A chest radiograph initially was interpreted as normal but on further review indicated mediastinal widening. A CT scan showed mediastinal lymphadenopathy, hemorrhagic mediastinitis, small bilateral pleural effusions, and a small pericardial effusion. Blood cultures grew *B. anthracis*. The patient is receiving ciprofloxacin, rifampin, and penicillin.

Florida

To date, the investigation in Florida has identified two confirmed inhalational cases. No new cases have been identified since the last report (1).

Clinical Presentation of Inhalational and Cutaneous Cases

Inhalational anthrax

To date, CDC has identified 10 patients with confirmed or suspected inhalational anthrax associated with bioterrorism. All but the most recent patients were postal workers (six), mail handlers or sorters (two), or a journalist who were known to or believed to have processed, handled, or received letters containing *B. anthracis* spores. The hospital employee with inhalational anthrax did not process mail but might have carried mail to other parts of the facility. Preliminary environmental testing of the patient's work area and home was negative for *B. anthracis*. The investigation is ongoing.

The median age of the 10 patients with inhalational anthrax was 56 years (range: 43–73 years); seven were men. The incubation period from the time of exposure to onset of symptoms when known (seven) was 7 days (range: 5–11 days).

The initial illness in these patients was characterized by fever (nine) and/or sweats/ chills (six) (Figure 2). Severe fatigue or malaise was present in eight and minimal or nonproductive cough in nine, including one with blood-tinged sputum. Eight patients reported chest discomfort or pleuritic pain. Abdominal pain or nausea or vomiting occurred in five, and five reported chest heaviness. Other symptoms included shortness of

breath (seven), headache (five), myalgias (four), and sore throat (two).

On initial presentation, total WBC count was normal or slightly elevated (7.5-13.3 x 103/cumm); however, elevation in the percentage of neutrophils or band forms was frequently noted. None of the patients had a low WBC count or lymphocytosis when initially evaluated. Chest radiograph was abnormal in all patients, but in two an initial reading was interpreted as within normal limits. Mediastinal changes including mediastinal widening, paratracheal fullness, hilar fullness, and mediastinal lymphadenopathy were noted in all eight patients who had CT scans. Mediastinal widening may be subtle, and careful review of the chest radiograph by a radiologist may be necessary. Pleural effusions were present in seven patients and were a feature of the two patients who did not have mediastinal changes on chest radiograph or did not have a CT scan. Pleural effusions often were large and hemorrhagic, reaccumulated, and required repeated thoracentesis or chest tubes. Pulmonary infiltrates were observed in four patients and were multilobar in three. Blood cultures grew B. anthracis in seven patients and in all who had not received antimicrobials. Diagnosis in the patients with negative cultures was confirmed by bronchial or pleural biopsy and specific IHC staining, by PCR of material from a sterile site, or by a fourfold rise in IgG to the protective antigen.

To date, six of 10 patients with inhalational anthrax have survived. Among those whose condition was recognized early, all remain alive and two have been discharged from the hospital. Prompt recognition of the early features of inhalational anthrax is important in settings of known or suspected exposure.

Cutaneous anthrax

Eleven patients with cutaneous anthrax have been identified in the current outbreak. Patients with cutaneous anthrax were mail handlers or sorters (four), employees of or visitors to media companies (six), and one bookkeeper. The mean incubation period for cutaneous anthrax was 5 days (range: 1–10 days) based on estimates from the postmark of letters and assumptions of dates of exposures with known positive letters or suspect letters (Figure 3).

Lesions occurred on the forearm, neck, chest, and fingers (two). Lesions were painless but accompanied by a tingling sensation or pruritis. Diagnosis was established by biopsy or culture.

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FIGURE 2. Clinical evaluation of persons with possible inhalational anthrax

History of exposure, or occupational/environmental risk with 2-5 day illness of: Symptoms Fever with or without chills Sweats, often drenching Fatique, malaise Cough (usually nonproductive), shortness of breath Chest discomfort, pleuritic pain Nausea, vomiting, diarrhea, abdominal pain Headache, myalgias Sore throat Signs Fever YES Observe closely Initial evaluation · Provide antimicrobial · Obtain white blood cell count (WBC), chest prophylaxis if exposure radiograph (CR), and blood cultures is confirmed (1) WBC: normal to elevated, neutrophilia with bands CR: · Mediastinal widening, · Pleural effusion, · Pulmonary infiltrate Consider chest computerized tomography (CT) if CR is normal Consider rapid diagnostic testing for influenza · Notify public health authorities WBC, CR, CT within Either WBC, CR, CT normal limits and abnormal or patient patient mildly ill moderately/severely ill Observe closely for Begin antimicrobial therapy (2) development of new If pleural effusion present, obtain fluid for gram symptoms stain and culture, polymerase chain reaction, Await blood cultures and cell block for immunohistochemistry* Initiate or continue If meningeal signs or altered mental status present,

- * Available through CDC or LRN. Cell block obtained by centrifugation of pleural fluid.
- ¹ Serologic testing available at CDC may be an additional diagnostic technique.

References

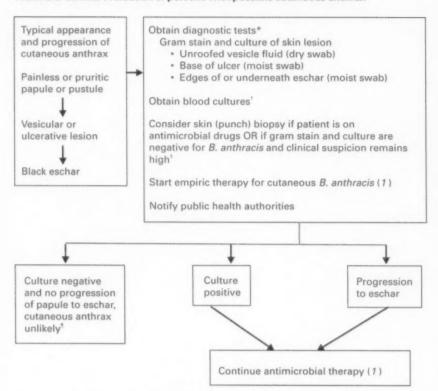
prophylaxis (1)

 CDC. Update: investigation of anthrax associated with intentional exposure and interim public health guidelines, October 2001. MMWR 2001;50:889–93.

perform lumbar puncture Other diagnostic tests'

CDC. Update: investigation of bioterrorism-related anthrax and interim guidelines for exposure management and antimicrobial therapy, October 2001. MMWR 2001;50:909–19.

FIGURE 3. Clinical evaluation of persons with possible cutaneous anthrax



 Serologic testing available at CDC may be an additional diagnostic technique for confirmation of cases of cutaneous anthrax.

¹ If blood cultures are positive for *B. anthracis*, treat with antimicrobials as for inhalational anthrax (1).

¹ Punch biopsy should be submitted in formalin to CDC. Polymerase chain reaction can also be done on formalin-fixed specimen. Gram stain and culture are frequently negative for *B. anthracis* after initiation of antimicrobials.

⁵ Continued antimicrobial prophylaxis for inhalational anthrax for 60 days if aerosol exposure to *B. anthracis* is known or suspected (2).

Reference

- CDC. Update: investigation of bioterrorism-related anthrax and interim guidelines for exposure management and antimicrobial therapy, October 2001, MMWR 2001:50:909-19.
- CDC. Update: investigation of anthrax associated with intentional exposure and interim public health guidelines, October 2001. MMWR 2001;50:889–93.

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Editorial Note: Since the last report (1), six new anthrax cases have been reported. Three of these cases have occupational exposures similar to previously reported cases (1). A fourth case occurred in a mail handler at a facility not previously linked to cases but that receives mail from a facility at which cases have occurred previously. Two new cases have no discernable epidemiologic link with anthrax cases previously reported or sites that are associated with known cases. These new cases suggest that anthrax exposure has occurred or is continuing to occur through means that cannot be ascribed to known contaminated letters or the paths these letters took through the mail service. The public health response to these new anthrax cases will evolve based on ongoing epidemiologic and criminal investigations.

Because exposures are being intentionally perpetrated, public health authorities must be vigilant for the appearance of new cases in previously unaffected populations. Prompt data sharing between law enforcement and public health authorities is essential.

Since September 11, 2001, state and local health departments have been responding to many reports of potential bioterrorist threats including letters containing powder, suspicious packages, and potential dispersal devices. During September 11–October 17, 40 state and territorial health officials who responded to a CDC telephone survey estimated that 7,000 reports had been received at their health departments, approximately 4,800 required phone follow-up, and 1,050 reports led to testing of suspicious materials at a public health laboratory (CDC, unpublished data, 2001). In comparison, the number of anthrax threats reported to federal authorities during 1996–2000 did not exceed 180 reported threats per year (Federal Bureau of Investigation, unpublished data, 2001). Therefore, although only four areas have identified cases of bioterrorism-associated anthrax, health departments throughout the nation are responding to public concerns, bioterrorism hoaxes, and threats.

CDC is working with state and local health departments and the U.S. Postal Service to develop standardized guidelines for identifying populations that should receive antimicrobial prophylaxis for prevention of inhalational anthrax. Current challenges include identifying factors that promote the aeresolization of *B. anthracis* in mail-handling facilities and assessing the risk for anthrax in environments contaminated with *B. anthracis* spores. Safe levels of *B. anthracis* spore contamination in occupational settings must be defined to determine the need for clean-up of contaminated facilities. The current antimicrobial prophylaxis recommendations address the prevention of inhalational anthrax, but CDC also is evaluating measures to prevent cutaneous anthrax.

Postexposure prophylaxis with a recommended antimicrobial agent for the prescribed period of time can prevent inhalational anthrax. In the case of a known contaminated letter sent to the office of a U.S. Senator, antimicrobial prophylaxis was administered to persons from the area of exposure and first-responders to the incident (1). To date, there have been no cases of anthrax, even among those who had the greatest exposure. Antimicrobial prophylaxis had been recommended for the U.S. State Department mail handler with anthrax, but the worker had not started treatment before the onset of illness. Public health response must include prompt initiation of prophylaxis for exposed persons and systems to promote adherence to a full 60-day regimen.

Previous guidelines recommended ciprofloxacin for antimicrobial prophylaxis until antimicrobial susceptibility test data was available (3). Isolates involved in the current bioterrorism attacks have been susceptible to ciprofloxacin, doxycycline, and several other antimicrobial agents. Considerations for choosing an antimicrobial agent include effectiveness, resistance, side effects, and cost. No evidence demonstrates that ciprofloxacin is more or less effective than doxycycline for antimicrobial prophylaxis to B. anthracis. Widespread use of any antimicrobial will promote resistance. Many common pathogens are already resistant to tetracyclines such as doxycycline. However, fluoroquinolone resistance is not yet common in these same organisms. To preserve the effectiveness of fluoroquinolone against other infections, use of doxycycline for prevention of B. anthracis infection among populations at risk may be preferable. However, the selection of the antimicrobial agent for an individual patient should be based on side-effect profiles, history of reactions, and the clinical setting.

CDC and state and local public health agencies continue to mobilize epidemiologic, laboratory, and other staff to identify and investigate acts of bioterrorism. Cases of bioterrorism-associated anthrax continue to occur and new risk populations may be identified. Until the cause of these acts are removed, public health authorities and clinicians should remain alert for cases of anthrax.

References

- CDC. Update: investigation of bioterrorism-related anthrax and interim guidelines for exposure management and antimicrobial therapy. MMWR 2001;50:909–19.
- CDC. Update: investigation of anthrax associated with intentional exposure and interim public health guidelines. MMWR 2001;50:889-93.
- Inglesby TV, Henderson DA, Bartlett JG, et al. Anthrax as a biological weapon: medical and public health management. JAMA 1999;281:1735–45.

Major Cardiovascular Disease (CVD) During 1997–1999 and Major CVD Hospital Discharge Rates in 1997 Among Women with Diabetes — United States

Cardiovascular disease (CVD) is the leading cause of death among all women (1) and the risk for death from CVD among women with diabetes is two to four times higher than that for women without diabetes (2). The excess risk for death as the result of CVD among persons with diabetes is better understood than the excess risk for CVD morbidity (2). To estimate national CVD prevalence and CVD hospital use among women with diabetes, CDC and the Agency for Health Care Research and Quality (AHRQ) analyzed data from the 1997–1999 National Health Interview Survey (NHIS) and the 1997 Nationwide Inpatient Sample (NIS). Findings indicate that the age-adjusted prevalence of major CVD for women with diabetes is twice that for women without diabetes and that the age-adjusted major CVD hospital discharge rate for women with diabetes is almost four times the rate for women without diabetes. These findings underscore the need to reduce risk factors associated with CVD among all women with diabetes through focused public health and clinical efforts.

The prevalence of CVD among women aged ≥18 years by diabetes status was obtained from a 3-year average of the estimates from the 1997–1999 NHIS, an ongoing nationally representative survey providing information on the health of the noninstitutionalized U.S. civilian population. Respondents were asked whether they had

Major Cardiovascular Disease - Continued

ever been told by a doctor or other health-care provider that they had hypertension, coronary heart disease, angina, heart attack, other kinds of heart conditions or heart disease, stroke, or diabetes. Major CVD was defined as one or more positive responses to the six CVD condition questions, and diabetes was defined as a positive response to the diabetes question.

The number of major CVD hospital discharges was estimated from the 1997 NIS, a stratified probability sample of hospitals in 22 states. Discharges from the 22 states represented approximately 60% of all discharges in the United States. Sample data were weighted using the American Hospital Association Annual Survey of Hospitals to approximate discharges from all U.S. acute-care community hospitals. Analysis was restricted to major CVD discharges (e.g., ischemic heart disease, hypertensive disease, rheumatic heart disease, and stroke) having an International Classification of Disease, Ninth Revision, Clinical Modification (ICD-9-CM) code of 390-448 as the first-listed diagnosis. Diabetes-related discharges were identified as those with an ICD-9 code of 250 as a secondary diagnosis. Major CVD discharge rates were calculated for the U.S. female population aged ≥18 years with and without diabetes by dividing the number of major CVD discharges estimated from NIS by the estimated number of women with and without diabetes obtained using 1997 NHIS data. Rate ratios and rate differences were calculated for both major CVD prevalence and hospital discharge rates by comparing rates for women with diabetes with those for women without diabetes. SUDAAN was used to calculate all estimates and their standard errors because of the complex sample designs of NIS and NHIS.

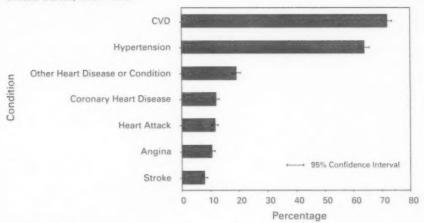
Major CVD rates were age-adjusted to the 2000 U.S. standard population (3). Trends were assessed by age for major CVD prevalence and for hospital discharge rates, rate ratios, and rate differences. The difference between age-adjusted major CVD prevalence and hospital discharge rates by diabetes status across all racial/ethnic categories was assessed using z-tests; a t-test in SUDAAN was used to assess the difference in age-adjusted major CVD prevalence and hospital discharge rates by race/ethnicity, comparing non-Hispanic whites, non-Hispanic blacks, and Hispanics. Other racial/ethnic groups were not included because sample size was too small for meaningful analysis.

During 1997–1999, 72% (95% confidence interval [CI]= \pm 1.8%) of all women with diabetes self-reported having major CVD (Figure 1). The most common CVD condition was hypertension (64%; 95% Cl= \pm 1.8%) followed by other heart disease or conditions (19%; 95% Cl= \pm 1.5%), coronary heart disease (12%; 95% Cl= \pm 1.2%), heart attack (11%; 95% Cl= \pm 1.3%), angina (10%; 95% Cl= \pm 1.1%), and stroke (8%; 95% Cl= \pm 1.0%) (Figure 1).

The prevalence of major CVD increased with age for women with diabetes from 40.5% (95% Cl= \pm 4.9%) for women aged 18–44 years to 85.1% (95% Cl= \pm 3.0%) for women aged \geq 75 years (p<0.0001) (Table 1). The age-adjusted prevalence of major CVD among women with diabetes was twice that of women without diabetes (p<0.0001) (Table 1). Age-adjusted major CVD prevalence for women with diabetes was higher for non-Hispanic blacks (65.2%; 95% Cl= \pm 5.3%) than for non-Hispanic whites (55.7%; 95% Cl= \pm 3.7%) (p=0.004) or Hispanics (56.6%; 95% Cl= \pm 6.5%) (p=0.05). The rate ratio of major CVD in women with diabetes relative to women without diabetes ranged from 3.0 (95% Cl= \pm 0.4) for women aged 18–44 years to 1.3 (95% Cl= \pm 0.1) for those aged \geq 75 years (p=0.09). Although rate differences were greatest among women aged 45–64 years and lowest among women aged \geq 75 years, no significant trend by age was found (p=0.27) (Table 1).

Major Cardiovascular Disease - Continued

FIGURE 1. Prevalence of major cardiovascular disease (CVD) and specific CVD conditions among women with diabetes — National Health Interview Survey, United States, 1997–1999*



* 3-year average.

During 1997, 772,346 of all major CVD hospital discharges (28%) had diabetes as a secondary diagnosis (Table 2). Hospital discharge rates for major CVD among women with diabetes increased from 22.9 per 1,000 (95% Cl=±4.5) for the youngest age group to 332.7 per 1,000 (95% Cl=±54.3) for the oldest age group (p=0.0004) (Table 2). The age-adjusted major CVD hospital discharge rate for women with diabetes was 3.8 times that of women without diabetes (Table 2). No significant difference was found among racial/ethnic groups for major CVD hospital discharge rates among women with diabetes. The rate ratio comparing major CVD hospital discharges in women with diabetes with those without diabetes decreased with age from 11.8 per 1,000 (95% Cl=±2.4) in the youngest age group to 2.4 per 1,000 (95% Cl=±0.4) in the oldest (p=0.04). Rate differences increased with age from 20.9 per 1,000 (95% Cl=±4.4) in the youngest to 196.5 per 1,000 (95% Cl=+55.2) in the oldest (p=0.02) (Table 2).

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Editorial Note: This report indicates that 72% of U.S. women with diabetes self-reported having major CVD. Major CVD prevalence is twice as common and major CVD hospitalizations are nearly four times as common among women with diabetes compared with women without diabetes. These findings are consistent with mortality studies documenting that women with diabetes are at much higher risk for death as a result of major CVD than women without diabetes (2).

Clinical trials indicate that antihypertensive treatment (4), aspirin use (5), and lipidlowering therapies (6) prevent or delay cardiovascular events. Epidemiologic evidence suggests that the risk for major CVD might be reduced through glycemic control (7) and the promotion of healthy lifestyles, including weight reduction/obesity prevention, TABLE 1. Number* and rate' of major cardiovascular disease (CVD) among women with and without diabetes, by age

Major Cardiovascular Disease — Continued

		Diabete	8	4	Vo diabe	ites	Rate		Rate	
Characteristic	No.	Rate	Rate (95% Cff)	No.	Rate	Rate (95% CI)	ratio	(95% CI)	difference	difference (95% CI)
Age group (yrs)										
18-44	367,750	40.5	(35.6-45.3)	7,121,007	13,3		3.0	(2.7-3.4)	27.2	(22.3-32.0)
45-CM	1,648,082	72.9	(70.2-75.6)	9,073,706	34.1		2.1	(2.0-2.2)	38.8	(36.0-41.6)
65-74	1,114,985	81.1	(78.1-84.0)	4,656,932	55.9	(54.3-57.4)	1.5	(1.4-1.5)	25.2	(21.9-28.5)
>75	865,584	85.1	(82.1-88.1)**	4,945,152	66.2	(64.7-67.6)**	1.3	(1.2-1.3)"	18.9	(15.6-22.2)
Inadjusted total	3,996,401	71.8	(70.1-73.6)	25,796,797	26.9	(26.4-27.3)	2.7	(2.6-2.8)	45.0	(43.1-46.8)
Age-adjusted total ¹¹		57.4	(54.7-60.1)		27.6	(27.2-28.0)	2.1	(2.0-2.2)	29.8	(27.0-32.5)
Race/Ethnicity th										
White, non-Hispanic	2,681,463	55.7	(52.0-59.4)	20,052,115	27.1	(26.6-27.6)	2.1	(1.9-2.2)	28.6	(24.9-32.3)
Black, non-Hispanic	814,748	65.2	(60.0-70.5)	3,463,197	36.4	(35.2-37.6)	1.8	(1.6-2.0)	28.9	(23.5-34.3)
Hispanic	403 153	56.6	(50.2-63.1)	1671038	24.1	1230-2531	20	19 9-2 61	22 6	126 0 30 11

* Weighted sample size.

' Per 100 women.

1.3-year average.

** Confidence interval.

** p.c.0.0001;t-test for trend.

** p.c.3.0242; linear regression test for trend.

** p.c.3.242; linear regression test for trend.

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		Diabetes	98		No diabetes	98	Rate		Rate	
Characteristic	No.	Rate	(95% CI ¹)	No.	Rate	(95% CI)	ratio	(BB% CI)	difference	(95% CI)
Age group (yrs)										
18-44	19,735	22.9	(18.4-27.4)	103,948	1.9	(1.8-2.1)	11.8	(9.4-14.3)	20.9	(16.5-25.4)
45-64	201,816	93.1	(81.7-104.5)	389,690	14.9	(14.0-15.8)	6.2	(5.4-7.1)	78.2	(66.7-89.7)
65-74	248,357	158.9	(134.6-183.2)	483,587	56.9	(52.6-61.3)	2.8	(2.3-3.3)	102.0	(77.4-126.6)
>75	302,296	332.7	(278.4-387.0)*	1,030,783	136.2	(126.4-146.0)1	2.4	(2.0- 2.9)*		(141,3-251,7)
Unadjusted total**	772,346	140.4	(127.9-152.9)	2,017,788	21.0	(20.0- 22.0)	6.7	(6.0- 7.4)	- fee	(106.9-131.8)
Age-adjusted total ⁷⁷		81.2	(73.6-88.8)		21.6	(20.4-22.8)	3.8	(3.3- 4.2)	59.5	(51.9-67.2)
White one Menoric	443 808	65.2	1 67 4 73 01	1 340 419	18.9	15.6. 18.31	a	133 441	48.3	1 40 3 56 31
Black, non-Hispanic	109,559	69.7	(53.8-85.6)	169,399	21.7	(17.7-25.7)	3.2	(2.3-4.2)	48.0	
Hispanic	46.127	57.0	(37.0-77.0)	71,037	16.1	(10.7-21.5)	50,00		41.0	(20.2-61.7)

* Weighted sample size.

* Per 1,000 women.

* Confidence interval.

* p.-0,001; trend.

* p.-0,003; linear regression test for trend.

* p.-0,0348; linear regression test for trend.

* p.-0,0348; linear regression test for trend.

* Includes missing ages.

* Mage-adjusted to the 2000 U.S. standard population. Numbers for other racial/ethnic groups were too small for meaningful analysis.

Major Cardiovascular Disease - Continued

increased physical activity, smoking cessation/prevention, and improved diet (7). Primary prevention of diabetes, a risk factor for major CVD, also can prevent major CVD. The Diabetes Prevention Program, a clinical trial examining the effect of intensive lifestyle intervention on the occurrence of type 2 diabetes in high-risk populations, concluded that improved diet, weight loss, and increased physical activity prevented or delayed the onset of diabetes among adults with impaired glucose tolerance (8).

Despite the efficacy of prevention strategies for major CVD, a large proportion of persons with diabetes have uncontrolled blood pressure (9), dyslipidemia (9), and hyperglycemia (9) and do not take aspirin (5). Additional research is needed to learn how to improve the process and outcomes of care among persons with diabetes. A concerted effort among health-care providers, public health officials, members of community-based organizations, and patients and their families will be necessary to reduce major CVD among persons with diabetes.

The high rate of major CVD among women with diabetes of all ages indicates that strategies for CVD risk reduction should be offered to all women with diabetes. Rate differences in hospital discharges increased with age, indicating that the effects of successful CVD prevention efforts should increase with age for women with diabetes.

The findings in this study are subject to at least six limitations. First, NHIS data on history of diabetes and major CVD are self-reported; however, studies comparing selfreported with physician-reported medical history data have found no difference in the prevalence of diabetes, and self-reported prevalence rates for CVD and hypertension were only slightly higher than physician-reported rates (10). Second, because NHIS excludes the institutionalized population, the number of persons with major CVD and diabetes is underestimated. Third, because NIS data represent hospital discharges and not individual persons, patients with multiple CVD hospitalizations within 1 year were counted multiple times; this might have resulted in an overestimation of hospital discharge rates. Fourth, by not including data from long-term and federal hospitals, NIS underestimates major CVD hospitalizations. Fifth, race/ethnicity is missing for approximately 20% of hospital discharges in NIS; four states that contribute to NIS provided no information on race/ethnicity, and one state provided race/ethnicity for approximately 25% of discharges. Therefore, race/ethnicity rates might be underreported and might be biased if disease patterns vary differentially across the reporting and nonreporting states. Finally, because the NIS sample comprised only 22 states, these data might be biased and might differ from estimates of the National Hospital Discharge Sample. However, in 1997, both data sources produced similar estimates of discharges with diabetes as the primary diagnosis (AHRQ, unpublished data, 2000).

CDC has published *Diabetes and Women's Health Across the Life Stages: A Public Health Perspective* that addresses the need for more research to gain a better understanding of the excess risk for major CVD among women with diabetes and to identify modifiable behavior and other determinants that can be used to develop effective interventions. The National Institutes of Health and CDC also have implemented the National Diabetes Education Program that includes public and private partners in the treatment and outcome of persons with diabetes; this program promotes early diagnosis to reduce morbidity and mortality associated with diabetes. CDC supports diabetes control programs in every state and, in 1998, initiated support for cardiovascular health promotion, disease prevention, and control programs. Since 1999, CDC has supported REACH 2010 (Racial and Ethnic Approaches to Community Health) to eliminate racial/ethnic disparities in numerous health areas, including diabetes and CVD. Additional information on diabetes is available at http://www.cdc.gov/Diabetes.

Major Cardiovascular Disease — Continued

References

- Hoyert DL, Kochanek KD, Murphy SL. Deaths: final data for 1997. National vital statistics reports; vol 47, no. 19. Hyattsville, Maryland: US Department of Health and Human Services, CDC, National Center for Health Statistics, 1999.
- Geiss LS, Herman WH, Smith PJ. Mortality in non-insulin-dependent diabetes. In: Harris MI, Cowie CC, Stern MP, Boyko EJ, Reiber GE, Bennett PH, eds. Diabetes in America. 2nd ed. Washington, DC: US Department of Health and Human Services, National Institutes of Health, publication no. 95-1468; 1995.
- Anderson RN, Rosenberg HM. Age standardization of death rates: implementation of the year 2000 standard. National vital statistics reports; vol 47, no. 3, Atlanta, Georgia: US Department of Health and Human Services, Public Health Service, CDC, National Center for Health Statistics, 1998.
- United Kingdom Prospective Diabetes 5 dy Group. Tight blood pressure control and risk of macrovascular and microvascular complications in type 2 diabetes: UKPDS 38. BMJ 1998;317:703–13.
- Rolka DB, Fagot-Campagna A, Venka arrayan KM. Aspirin use among adults with diabetes: estimates from the Third National Health and Nutrition Examination Survey. Diabetes Care 2001;24:197–201.
- Pyorala K, Pedersen TR, Kjekshus J, Faergeman O, Olsson AG, Thorgeirsson G. Cholesterol lowering with simvastatin improves prognosis of diabetic patients with coronary heart disease. Subgroup analysis of the Scandinavian Simvastatin Survival Study (4S) Group. Diabetes Care 1997;20:614–20.
- Haffner SM. Management of dyslipidemia in adults with diabetes. Diabetes Care 1998;21:160–78.
- National Institute of Diabetes and Digestive and Kidney Diseases. Diet and kidney diseases. [Press release]. August 8, 2001. Available at http://www.niddk.nih.gov/welcome/releases/8 8 01.htm>. Accessed August 10, 2001.
- Harris MI. Health care and health status and outcomes for patients with type 2 diabetes. Diabetes Care 2000;23:754–8.
- Kehoe R, Wu S-Y, Leske MC, Chylack LT Jr. Comparing self-reported and physicianreported medical history. Am J Epidemiol 1994;139:813–8.

Hospital Discharge Rates for Nontraumatic Lower Extremity Amputation by Diabetes Status — United States, 1997

Lower extremity amputation (LEA) is a costly and disabling procedure that disproportionately affects persons with diabetes (1,2). One of the national health objectives for 2000 was to reduce the LEA rate from a 1991 baseline of approximately eight per 1,000 persons with diabetes to a target of approximately five per 1,000 persons with diabetes. Review of 1996 data indicated an LEA rate of approximately 11. To estimate the national rates of hospital discharges for LEA among persons with and without diabetes and to assess the excess risk for LEA among persons with diabetes, CDC and the Agency for Healthcare Research and Quality (AHRQ) analyzed data from the 1997 Nationwide Inpatient Sample (NIS) and the 1997 National Health Interview Survey (NHIS). This report summarizes the findings of the analysis, which indicated that the age-adjusted rates of hospital discharges among persons with LEA who had diabetes were 28 times that of those without diabetes. This higher rate underscores the need to increase efforts to prevent risk factors (e.g., peripheral vascular disease, neuropathy, and infection) that result in LEA among persons with diabetes.

Nontraumatic Lower Extremity Amputation — Continued

Hospital discharges were estimated from NIS, a stratified probability sample of hospitals in 22 states. Discharges from these states represented approximately 60% of all discharges in the United States. Sample data were weighted using the American Hospital Association Annual Survey of Hospitals to approximate discharges from all U.S. acute-care community hospitals. LEA discharges were defined using International Classification of Disease, Ninth Revision, Clinical Modification (ICD-9-CM) codes 84.10-84.19 (traumatic LEA codes 895-897 were excluded). Diabetes-related LEA discharges were identified as discharges that included ICD-9-CM code 250 as one of the listed discharge diagnoses. LEA hospital discharge rates were calculated for populations with and without diabetes. Estimates of the number of persons with and without diabetes were obtained from the 1997 NHIS, an ongoing, nationally representative survey providing information about the health of the noninstitutionalized U.S. civilian population (3). SUDAAN was used to calculate estimates and 95% confidence intervals (CIs) of NIS and NHIS data. The 2000 U.S. standard population was used to adjust LEA rates by age. The rate ratio was calculated by dividing the LEA rate for persons with diabetes by the rate for persons without diabetes. The rate difference was defined as the difference in LEA rates between the two populations. The significance of trends by age was assessed for LEA rates, rate ratios, and rate differences, and t-tests in SUDAAN were used to determine the significance of the difference in mean age by diabetes status and the ageadjusted rate differences by sex and race. Z-tests were used to assess the difference between age-adjusted rates by diabetes status among all sex and race groups.

In 1997, 131,218 hospital discharges had an LEA discharge diagnosis code; 87,720 (67%) of these were related to diabetes (Table 1). Among persons with diabetes, 66.7% of LEA hospitalizations were paid by Medicare and an additional 8.1% were paid by Medicaid. Among persons with diabetes, approximately 52% of amputations occurred at or below the foot, and among persons without diabetes, approximately 70% occurred between the ankle and the knee or higher. Patients with diabetes-related LEA hospital discharges had a mean age of 66 years (95% Cl=±0.3 years), and the mean age of LEA discharges not related to diabetes was 71 years (95% Cl=±0.7 years) (p<0.0001). LEA rates increased with age in both populations, but rates were higher in the population with diabetes. LEA rate ratios ranged from 149 (95% Cl=116–182) to nine (95% Cl=7–10) for persons aged ≤44 years and ≥75 years, respectively. Rate differences ranged from 3.4 to

13.8 per 1,000 persons in those aged ≤44 years to ≥75 years, respectively.

The age-adjusted LEA rate for persons with diabetes (5.5 per 1,000 persons with diabetes) was 28 (95% CI=24–31) times that of persons without diabetes (0.2 per 1,000 persons without diabetes). Regardless of diabetes status, these rates were higher for men than women (p<0.0001) and higher for non-Hispanic blacks than Hispanics or non-Hispanic whites (p<0.05) (Figure 1). Age-adjusted LEA rates were much higher for persons with diabetes for both sexes and all racial/ethnic populations (p<0.0001).

Reported by: Center for Organization and Delivery Studies, Agency for Healthcare Research and Quality, Rockville, Maryland. Epidemiology and Statistics Br, Div of Diabetes Translation, National Center for Chronic Disease Prevention and Health Promotion, CDC.

Editorial Note: The findings in this report indicate that LEAs occur at a much higher rate among persons with diabetes and that diabetes causes approximately 67% of LEAs. Among persons with diabetes, LEA rates were highest among men, non-Hispanic blacks, and the elderly. These findings indicate that LEAs might increase as the U.S. population ages and as the prevalence of diabetes increases. Because approximately 75% of LEA

TABLE 1. Number of nontraumatic lower extremity amputation (LEA) hospital discharges, rate * of LEA for persons with and without diabetes, and LEA rate difference - National Inpatient Sample and National Health Interview

Nontraumatic Lower Extremity Amputation - Continued

		Diabete	S	No	diabet	Sa	Ra	te difference
Characteristic	No.	Rate	(95% CI')	No.	Rate	(12 % S6)	No.	(12 % S6)
ge group (yrs)								
0-44	5,844	3.4	(2.8-4.0)	4,121	0.02	(0.02-0.03)	3.4	(2.8-3.9)
45-64	31,352	7.5	(6.7-8.3)	7,916	0.16	(0.14-0.17)	7.3	(6.5-8.2)
65-74	25,362	8.6	(8.5-11.1)	9,670	0.62	(0.57-0.68)	9.2	(7.9-10.5)
≥75	25,162	15.6	(13.5-17.7)	21,767	1.78	(1.64-1.92)*	13.8	(11.7-16.0)*
otal	87,720	8.7	(8.0-9.4)	43,498**	0.17	(0.16 - 0.18)	8.5	(7.8-9.2)
ae-adjusted"		5.4	(4.9-6.0)		0.20	(0.19-0.21)	5.3	(4.7-5.8)

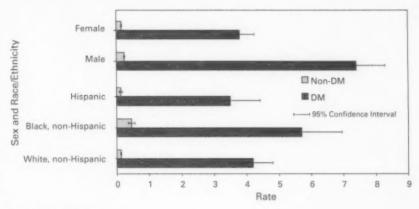
* Per 1,000 persons.

p<0.0001; t-test for trend. p=0.01; tinear regression test for trend. ** Includes missing ages.

" Age-adjusted to the 2000 U.S. standard population.

Nontraumatic Lower Extremity Amputation — Continued

FIGURE 1. Age-adjusted lower extremity amputation hospital discharge rates* for persons with diabetes mellitus (DM) and without diabetes mellitus (non-DM), by sex¹ and race/ethnicity⁵ — National Inpatient Sample and National Health Interview Survey, United States, 1997



*Per 1,000 persons.

1 p<0.0001; t-test for difference.

bp<0.05; t-test for difference among non-Hispanic blacks, non-Hispanic whites, and Hispanics. Numbers for other racial/ethnic groups were too small for meaningful analysis.

hospitalizations are paid by Medicare or Medicaid, the increase in prevalence will place a large financial burden on these public health systems.

Among persons with diabetes, LEAs result from the single or combined effects of peripheral vascular disease, peripheral neuropathy, and infection (1,4). Foot deformities and ulcers occurring as a consequence of neuropathy and/or peripheral vascular disease, minor trauma, and poor foot care also might contribute to LEAs (1,5).

The findings in this study are subject to at least five limitations. First, because NIS data represent hospital discharges and not individual persons, patients with multiple amputations within 1 year were counted multiple times; this might have resulted in an overestimation of hospital discharge rates. Second, because NIS data do not include LEAs that occurred in federal hospitals and outpatient settings, the analysis underestimates the total number of LEA discharges that occurred nationally. Third, because NHIS is representative of the noninstitutionalized civilian population, the total population with or without diabetes was underestimated. Fourth, race/ethnicity data are missing for approximately 20% of the hospital discharges in NIS data; four states contributing to NIS provided no race/ethnicity data and one state provided race/ethnicity information for approximately 25% of discharges. Therefore, race/ethnicity-specific rates are underreported and may be biased if race/ethnicity disease patterns vary across reporting and nonreporting states. Finally, because the NIS sample was constructed from only 22 states, these data might be biased and might differ from estimates of the National Hospital Discharge Sample (NHDS), However, in 1997, both data sources produced similar estimates of discharges with diabetes as the primary diagnosis (AHRQ, unpublished data, 2000).

Nontraumatic Lower Extremity Amputation - Continued

Serious foot conditions or LEA can be decreased by 44%–85% in persons with diabetes (5). Proper footware can lower abnormal pressure and protect the foot from calluses and ulcers, precursors of LEA (6). Education intervention, multidisciplinary care, and insurance coverage for therapeutic shoes are effective in reducing diabetes-related LEA (2). Interventions also include early detection of feet at risk through regular foot examination, knowledge of foot hygiene, nonweight-bearing exercise, and provider education on screening examinations for high-risk foot conditions (6,7). Good glycemic control can reduce the development of neuropathy, a high-risk condition for LEA (8,9).

Because no nationally representative data on lower extremity disease and its risk factors exist, in 1999, CDC and the National Heart, Lung, and Blood Institute of the National Institutes of Health added to the National Health and Nutrition Examination Survey a lower extremity disease examination component for peripheral vascular disease, peripheral neuropathy, and foot deformities, ulcers, and amputations. This component will allow national estimates of the extent of lower extremity disease and identification of its risk factors. It also will increase an understanding of racial/ethnic differences in lower extremity disease and provide information to clinicians and public health providers to develop preventive care and community-based interventions. Materials designed to make good foot care an essential part of diabetes care among health-care providers and persons with diabetes are available at https://ndep.nih.gov/materials/pubs/feet/feet.htm.

References

- Reiber GE, Boyko EJ, Smith DG. Lower extremity foot ulcers and amputations in diabetes. In: Harris MI, Cowie CC, Stern MP, Boyko EJ, Reiber GE, Bennett PH, eds. Diabetes in America. 2nd ed. Bethesda, Maryland: National Diabetes Data Group of the National Institute of Diabetes and Digestive and Kidney Diseases, National Institutes of Health, 1995:409– 28. Available at http://diabetes-in-america.s-3.com/. Accessed October 2001.
- Ollendorf DA, Kotsanos JG, Wishner WJ, et al. Potential economic benefits of lower extremity amputation prevention strategies in diabetes. Diabetes Care 1998;21:1240–5.
- National Center for Health Statistics. Data file documentation, National Health Interview Survey, 1997 [machine-readable data file and documentation]. Hyattsville, Maryland: US Department of Health and Human Services, Public Health Service, National Center for Health Statistics, 1997.
- CDC. The prevention and treatment of complications of diabetes mellitus: a guide for primary care practitioners. Atlanta, Georgia: US Department of Health and Human Services, Public Health Service, CDC, 1991.
- Bild DE, Selby JV, Sinnock P, Browner WS, Braveman P, Showstack JA. Lower extremity amputation in people with diabetes: epidemiology and prevention. Diabetes Care 1989;12:24–31.
- Mayfield JA, Reiber GE, Sanders LJ, Janisse D, Pogach LM. Preventive foot care in people with diabetes. Diabetes Care 1998;21:2161–77.
- American Diabetes Association. Preventive foot care in people with diabetes. Diabetes Care 2001;24(suppl 1):S56–S57.
- Ohkubo Y, Kishikawa H, Araki E, et al. Intensive insulin therapy prevents the progression of diabetic microvascular complications in Japanese patients with non-insulin-dependent diabetes mellitus: a randomized prospective 6-year study. Diabetes Res Clin Pract 1995;28:103–17.
- Diabetes Control and Complications Trial Research Group. The effect of intensive treatment of diabetes on the development and progression of long-term complications in insulin-dependent diabetes mellitus. N Engl J Med 1993;329:977-86.

Weekly Update: West Nile Virus Activity — United States, October 24–30, 2001

The following report summarizes West Nile virus (WNV) surveillance data reported to CDC through ArboNET and verified by states and other jurisdictions as of October 30, 2001.

During the week of October 24–30, no human cases of WNV encephalitis or meningitis were reported. During the same period, WNV infections were reported in 200 crows, 43 other birds, and eight horses. A total of 11 WNV-positive mosquito pools were reported in five states (Georgia, Kentucky, New Jersey, Ohio, and Virginia).

During 2001, a total of 37 human cases of WNV encephalitis or meningitis have been reported in Florida (ten), Maryland (six), New Jersey (six), New York (six), Connecticut (five), Pennsylvania (three), and Georgia (one); one death occurred in Georgia. Among these 37 cases, 20 (54%) were in men; the median age was 69 years (range: 36–81 years); and dates of illness onset ranged from July 13 to October 7. A total of 3,996 crows and 1,437 other birds with WNV infection were reported from 25 states and the District of Columbia (Figure 1); 159 WNV infections in other animals (all horses) were reported from 13 states (Alabama, Connecticut, Florida, Georgia, Kentucky, Louisiana, Massachusetts, Mississippi, New York, North Carolina, Pennsylvania, Tennessee, and Virginia); and 736 WNV-positive mosquito pools were reported from 15 states (Connecticut, Florida, Georgia, Illinois, Kentucky, Maryland, Massachusetts, Michigan, New Hampshire, New Jersey, New York, Ohio, Pennsylvania, Rhode Island, and Virginia).

Additional information about WNV activity is available at http://cindi.usgs.gov/hazard/event/west_nile/west_nile.htm>.

FIGURE 1. Areas reporting West Nile virus (WNV) activity — United States, 2001*



^{*} As of October 30, 2001.

^{&#}x27; Mississippi reported WNV infection in a horse but no birds.

Notice to Readers

Updated Recommendations for Antimicrobial Prophylaxis Among Asymptomatic Pregnant Women After Exposure to Bacillus anthracis

The antimicrobial of choice for initial prophylactic therapy among asymptomatic pregnant women exposed to *Bacillus anthracis* is ciprofloxacin, 500 mg twice a day for 60 days. In instances in which the specific *B. anthracis* strain has been shown to be penicillinsensitive, prophylactic therapy with amoxicillin, 500 mg three times a day for 60 days, may be considered. Isolates of *B. anthracis* implicated in the current bioterrorist attacks are susceptible to penicillin in laboratory tests, but may contain penicillinase activity (2). Pencillins are not recommended for treatment of anthrax, where such penicillinase activity may decrease their effectiveness. However, penicillinase are likely to be effective for preventing anthrax, a setting where relatively few organisms are present. Doxycycline should be used with caution in asymptomatic pregnant women and only when contraindications are indicated to the use of other appropriate antimicrobial drugs.

Pregnant women are likely to be among the increasing number of persons receiving antimicrobial prophylaxis for exposure to *B. anthracis*. Clinicians, public health officials, and women who are candidates for treatment should weigh the possible risks and benefits to the mother and fetus when choosing an antimicrobial for postexposure anthrax prophylaxis. Women who become pregnant while taking antimicrobial prophylaxis should continue the medication and consult a health-care provider or public health official to discuss these issues.

No formal clinical studies of ciprofloxacin have been performed during pregnancy. Based on limited human information, ciprofloxacin use during pregnancy is unlikely to be associated with a high risk for structural malformations in fetal development. Data on ciprofloxacin use during pregnancy from the Teratogen Information System indicate that therapeutic doses during pregnancy are unlikely to pose a substantial teratogenic risk, but data are insufficient to determine that there is no risk (1). Doxycycline is a tetracycline antimicrobial. Potential dangers of tetracyclines to fetal development include risk for dental staining of the primary teeth and concern about possible depressed bone growth and defective dental enamel. Rarely, hepatic necrosis has been reported in pregnant women using tetracyclines. Penicillins generally are considered safe for use during pregnancy and are not associated with an increased risk for fetal malformation. Pregnant women should be advised that congenital malformations occur in approximately 2%–3% of births, even in the absence of known teratogenic exposure.

Additional information about the treatment of anthrax infection is available at http://www.cdc.gov/mmwr/preview/mmwrhtml/mm5042a1.htm.

Reference

- Friedman JM, Polifka JE. Teratogenic effects of drugs: a resource for clinicians (TERIS).
 Baltimore, Maryland: Johns Hopkins University Press, 2000:149–95.
- CDC. Update: investigation of bioterrorism-related anthrax and interim guidelines for exposure management and antimicrobial therapy, October 2001. MMWR 2001;50:909–19.

Notices to Readers — Continued Notice to Readers

Interim Recommendations for Protecting Workers from Exposure to *Bacillus anthracis* in Work Sites in which Mail is Handled or Processed

CDC has developed interim recommendations to assist personnel responsible for occupational health and safety in developing a comprehensive program to reduce potential cutaneous or inhalational exposures to *Bacillus anthracis* spores among workers in work sites in which mail is handled or processed. Such work sites include post offices, mail distribution/handling centers, bulk mail centers, air mail facilities, priority mail processing centers, public and private mail rooms, and other settings in which workers are responsible for handling and processing mail. The recommendations are based on the limited information available on methods to avoid infection and on the effectiveness of various prevention strategies. These recommendations will be updated as new informat on becomes available.

The recommendations are divided into the following hierarchical categories describing measures that should be implemented in distribution/handling centers to prevent potential exposures to *B. anthracis* spores:

- · Engineering controls to prevent or capture aerosolized spores
- Administrative controls to limit the number of persons potentially exposed to spores
- · Housekeeping controls to further reduce the spread of spores
- Personal protective equipment for workers to prevent cutaneous and inhalational exposure to spores

These control measures should be selected on the basis of an initial work site evaluation that focuses on determining which processes, operations, jobs, or tasks would be most likely to result in an exposure if a contaminated envelope or package enters the work site. The complete interim recommendations are available at http://www.bt.cdc.gov>.

Notice to Readers

National Diabetes Awareness Month — November 2001

November is National Diabetes Awareness Month. During 1998 in the United States, an estimated 15.7 million persons had diabetes (1). From 1990 to 2000, an increase of 49% occurred in the prevalence of diagnosed diabetes and gestational diabetes in U.S. adults (2); however, lifestyle changes, including weight control and regular physical activity can prevent or delay the onset of type 2 diabetes, even in high-risk persons (3).

During November, 59 state and territorial diabetes control programs, other partners, and CDC will highlight activities that increase awareness of the Initiative on Diabetes and Women's Health and of the need for persons with diabetes to receive influenza and pneumococcal vaccines. Persons with diabetes should receive pneumococcal and annual influenza vaccinations because they are more likely than persons without diabetes to die from complications of influenza and pneumonia (4). In 1997, only half of adults with diabetes received an annual influenza vaccination, and one third received a pneumococcal vaccine (5).

Notices to Readers - Continued

CDC, the American Diabetes Association, the American Public Health Association, and the Association of State and Territorial Health Officials cosponsor the Initiative on Diabetes and Women's Health, which has three phases: a report; the National Public Health Action Plan for Women and Diabetes; and a national conference. CDC's recently published report, Diabetes and Women's Health Across the Life Stages: A Public Health Perspective, is the first major publication to address the unique and serious impact diabetes has on women throughout life and to address the public health implications of these issues (6). The publication presents 1) trends in risk factors for diabetes and its complications during adolescence; 2) the increased risk for offspring to develop diabetes associated with intrauterine exposure to hyperglycemia; 3) the effect of menopause on health status; and 4) the increase in poverty and disability for older women.

Additional information about diabetes is available from CDC, telephone (877) 232-3422, e-mail diabetes@cdc.gov, and from http://www.cdc.gov/diabetes.

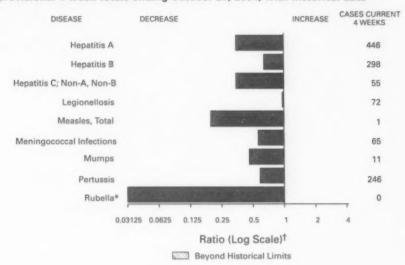
References

- CDC. National diabetes fact sheet: national estimates and general information on diabetes in the United States. Atlanta, Georgia: US Department of Health and Human Services, CDC, 1998.
- Mokdad AH, Bowman BA, Ford ES, et al. The continuing epidemics of obesity and diabetes in the United States. JAMA 2001;286:1195–200.
- Diabetes Prevention Program. National Institutes of Health. 2001 [1 screen]. Available at http://www.niddk.nih.gov/patient/dpp/dpp-q&a.htm. Accessed October 2001.
- Valdez R, Narayan KM, Geiss LS, Engelgau MM. Impact of diabetes mellitus on mortality associated with pneumonia and influenza among non-Hispanic black and white US adults. Am J Public Health 1999;89:1715–21.
- CDC. Influenza and pneumococcal vaccination rates among persons with diabetes mellitus—United States, 1997. MMWR 1999;48:961-7.
- Beckles GLA, Thompson-Reid PE, eds. Diabetes and women's health across the life stages: a public health perspective. Atlanta, Georgia: US Department of Health and Human Services, CDC, National Center for Chronic Disease Prevention and Health Promotion, Division of Diabetes Translation, 2001.

Erratum: Vol. 50, No. 42

In the article, "Update: Investigation of Bioterrorism-Related Anthrax and Interim Guidelines for Exposure Management and Antimicrobial Therapy, October 2001," on page 911 two dates were incorrect. The fourth sentence of the fourth paragraph should read "The patient was reported to CDC on October 15, and serologic testing at CDC was positive to B. anthracis." The first sentence of the sixth paragraph should read, "On October 18, the postal facility was closed; the New Jersey Department of Health and Senior Services recommended that postal workers at both postal facilities initiate antimicrobial prophylaxis pending further epidemiologic and environmental investigation."

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending October 27, 2001, with historical data



'No rubella cases were reported for the current 4-week period yielding a ratio for week 43 of

Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending October 27, 2001 (43rd Week)*

		Cum. 2001		Cum. 2001
Anthrax		11	Poliomyelitis, paralytic	
Brucellosis ¹		73	Psittacosis¹	15
Cholera		3	Q fever*	18
Cyclosporiasis	§ ¹	122	Rabies, human	1
Diphtheria		2	Rocky Mountain spotted fever (RMSF)	475
Ehrlichiosis:	human granulocytic (HGE)1	171	Rubella, congenital syndrome	-
	human monocytic (HME)1	71	Streptococcal disease, invasive, group A	2,938
Encephalitis:	California serogroup viral ¹	84	Streptococcal toxic-shock syndrome ¹	41
	eastern equine1	6	Syphilis, congenital [¶]	166
	St. Louis'	1	Tetanus	22
	western equine1		Toxic-shock syndrome	96
Hansen diseas		71	Trichinosis	22 96 21 90
	Imonary syndrome ¹	7	Tularemia*	90
	emic syndrome, postdiarrheal	121	Typhoid fever	227
HIV infection.		153	Yellowfever	
Plaque		2		

No reported cases.

*Incidence data for reporting year 2001 are provisional and cumulative (year-to-date).

Not notifiable in all states.

*Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV,

STD, and TB Prevention (NCHSTP). Last update September 25, 2001 Updated from reports to the Division of STD Prevention, NCHSTP.

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending October 27, 2001, and October 28, 2000 (43rd Week)*

	AII	ne	Chlam	wdia!	Comton	oridiosis	NET		coli O157:H7 PH	
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
Reporting Area	2001	2000	2001	2000	2001	2000	2001	2000	2001	2000
INITED STATES	29,580	31,406	579,265	573,902	2,403	2,621	2,477	3,923	1,959	3,209
NEW ENGLAND Maine N.H. /t. Mass. R.I.	1,129 36 31 13 602 78	1,639 28 28 29 1,049 75	18,705 893 1,104 506 7,747 2,422	19,167 1,192 909 439 8,148 2,216	109 17 14 30 44 4	127 20 21 26 33 3	206 25 32 13 111	341 26 31 32 154 18	208 26 26 8 106 11	354 28 34 33 161
Conn.	369	430	6,033	6,263		24	11	80	31	81
MID. ATLANTIC Jpstate N.Y. N.Y. City N.J. Pa.	6,710 731 3,385 1,389 1,206	6,816 664 3,695 1,345 1,112	62,871 11,368 24,324 8,936 18,243	53,734 1,961 21,927 8,890 20,956	221 89 73 7 52	330 105 152 16 57	184 142 11 31 N	390 258 21 111 N	165 121 10 34	274 60 15 111 88
E.N. CENTRAL Ohio nd. III. Mich. Wis.	2,238 430 264 992 413 139	3,042 430 282 1,568 601 161	91,419 20,222 12,732 22,501 24,980 10,984	98,533 25,947 11,017 27,604 20,443 13,522	911 140 72 1 163 535	869 239 57 109 84 380	664 174 72 144 81 193	968 240 109 180 130 309	450 137 39 128 69 77	687 204 83 145 103 152
W.N. CENTRAL Minn. lowa Mo. N. Dak. S. Dak. Nebr. Kans.	637 108 71 312 2 22 52 70	723 129 69 349 2 7 53 114	29,666 5,994 3,797 10,826 750 1,414 2,175 4,710	32,651 6,727 4,437 11,104 729 1,521 3,067 5,066	354 137 75 36 12 6 88	328 112 72 27 15 15 78 9	395 151 76 46 17 37 51	568 149 169 98 15 53 58 26	404 186 59 77 31 40	541 173 139 90 20 57 46 16
S. ATLANTIC Del. Md. D.C. Va. W. Va. N. C. S.C. Ga.	9,497 203 1,506 644 723 61 726 577 1,031 4,026	8,757 182 1,056 570 574 50 585 639 991 4,110	109,994 2,041 9,278 2,533 14,965 1,937 16,621 9,374 23,903 29,342	108,234 2,370 11,767 2,696 12,815 1,779 18,505 7,954 22,752 27,596	272 6 34 10 24 2 24 103 69	404 5 9 13 16 3 21 148 189	198 4 23 47 10 46 10 26 32	318 2 30 1 61 14 77 21 35	121 6 1 U 36 8 29 11 15	260 1 2 U 57 12 65 16 36 71
E.S. CENTRAL Ky. Tenn. Ala. Miss.	1,423 278 456 347 342	1,609 160 684 417 348	39,887 7,415 11,910 11,037 9,525	42,269 6,638 12,148 13,082 10,401	39 4 12 13 10	45 5 11 15 14	115 57 35 16 7	122 39 51 8 24	96 47 36 6 7	98 31 47 9
W.S. CENTRAL Ark. La. Okla. Tex.	3,141 159 665 186 2,131	3,333 158 554 294 2,327	86,710 6,043 14,403 8,591 57,673	86,903 5,547 15,261 7,826 58,269	32 6 7 12 7	146 11 10 17 108	84 12 4 26 42	213 54 14 18 127	86 25 24 37	262 37 44 16 165
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	1,073 14 17 3 231 103 437 90 178	1,211 12 19 9 294 126 386 113 252	33,131 1,542 1,542 676 7,022 4,738 11,974 1,512 4,125	32,048 1,154 1,513 662 8,809 4,202 10,589 1,851 3,268	193 30 21 6 34 22 7 69 4	156 10 21 5 65 16 10 25 4	248 16 63 5 83 13 22 31 15	382 30 64 17 147 19 44 48	127 1 52 9 22 42	281 38 9 105 16 35 68 10
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	3,732 395 154 3,112 16 55	4,276 379 113 3,670 15 98	106,882 11,264 6,122 84,121 2,180 3,195	100,363 10,740 5,499 79,092 2,062 2,970	272 43 44 181 1	216 U 16 200	383 109 61 192 4 17	621 195 126 257 29	302 62 57 176 1 6	452 196 107 135 3
Guam P.R. V.I. Amer. Samoa C.N.M.I.	10 934 2	13 1,052 31	2,140 53 U 103	422 U	Ü		N 1 U	N 6	0	U

Not notifiable. U: Unavailable. <: No reported cases. C.N.M.L.: Commonwealth of Northern Mariana Islands.
Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date). N: Not notifiable. cumulative (year-to-date).

cumulative (year-10-date). Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

Chlamydia refers to genital infections caused by C. trachomatis.

Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention. Last updated September 25, 2001.

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending October 27, 2001, and October 28, 2000 (43rd Week)*

	Gonorr	hea	Hepatit Non-A, N	is C; Ion-B	Legione	llosis	Listeriosis	Ly Disa	me ease
Reporting Area	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2001	Cum. 2000
UNITED STATES	267,135	294,333	2,683	2,634	820	914	374	10,536	14,235
NEW ENGLAND Maine N.H. Vt. Mass. R.I. Conn.	5,408 97 154 53 2,445 681 1,978	5,413 77 89 54 2,229 536 2,428	14 6 8	24 2 4 13 5	56 8 10 5 15 9	50 2 2 5 16 8 17	31 1 4 2 19 1	3,337 130 14 654 436 2,103	4,379 60 32 1,094 414 2,779
MID. ATLANTIC Jpstate N.Y. N.Y. City N.J. Pa.	31,792 6,932 10,238 5,856 8,766	32,139 6,094 9,554 6,005 10,486	1,310 48 1,214 48	589 33 517 39	161 56 16 8 82	251 73 41 20 117	57 25 8 10 14	5,301 2,990 2 927 1,382	7,522 3,241 170 2,346 1,765
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	49,109 11,199 5,603 13,874 14,464 3,969	58,968 15,875 5,185 17,360 14,753 5,795	148 5 1 13 129	197 11 19 167	226 109 19 65 33	238 99 32 28 42 37	50 13 8 1 21 7	577 105 20 21 1 430	744 56 22 34 23 609
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak.	12,758 1,964 997 6,741 33 228	14,783 2,637 1,048 7,266 60 253	603 9 582	484 5 2 466	46 9 8 19 1 3	54 7 13 24	15	337 279 30 23	361 267 30 45
Nebr. Kans.	710 2.085	1,217	3 9	4 7	5	4	1 4	3 2	3 15
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	68,261 1,212 5,030 2,292 8,948 13,895 6,293 13,168 16,869	76,885 1,418 8,128 2,163 8,673 539 15,135 7,091 14,746 18,992	96 16 9 19 6	92 12 3 3 14 14 2 3 39	170 11 32 7 20 N 8 10 9 73	165 8 60 5 31 N 13 4 6	11 11 5 5 5 11	740 49 474 10 111 11 37 5	995 167 583 5 133 28 43 7
E.S. CENTRAL Ky. Tenn. Ala. Miss.	25,780 2,967 8,031 8,515 6,267	30,500 2,942 9,738 10,130 7,690	169 8 57 4 100	393 31 85 9 268	50 11 25 12 2	32 17 10 3 2	19 5 8 6	53 22 22 22 8	47 11 28 5 3
W.S. CENTRAL Ark. La. Okla. Tex.	42,336 3,646 9,857 3,897 24,936	45,836 3,257 11,248 3,450 27,881	171 4 83 3 81	633 8 379 8 238	2 3	7 3 12	17 1 2 14	79 1 78	78 5 7
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	8,313 96 62 67 2,412 799 3,240 119 1,528	8,724 39 69 40 2,661 923 3,533 178 1,281	59 1 2 6 19 11 9 3	66 4 3 2 12 13 18 1	46 3 1 13 2 18 5 4	37 1 5 13 1 7	31 1 7 7 6 2 7	12 6 1 1 1 2	11 2 3 3 3 3
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	23,378 2,499 963 19,080 347 489	21,085 1,911 790 17,713 287 384	113 19 12 82	156 28 25 101	60 8 N 48	65 15 N 49	92 8 8 70	100 8 8 82 2 N	98 7 9 80 2 N
Guam P.R. V.I. Amer, Samoa C.N.M.I.	496 6 U	45 421 U	i Ü	3 1 U	2	1	:	Ň	NUU

N: Not notifiable. U: Unavailable. -: No reported cases.
* Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States,

						Salmon	ellosis ¹	
		laria		s, Animal	NET			ILIS
Reporting Area	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000
INITED STATES	955	1,237	5,559	5,961	30,139	32,734	24,463	27,634
NEW ENGLAND Maine N.H. 11. Mass. R.I. Conn.	64 4 2 1 27 7 23	65 6 1 2 30 8 18	620 60 20 56 223 56 203	701 116 21 53 231 50 230	2,076 158 154 71 1,154 113 426	1,910 108 123 100 1,100 121 358	1,963 137 139 63 1,054 154 416	1,945 88 130 96 1,106 132 393
MID. ATLANTIC Jpstate N.Y. V.Y. City V.J.	256 58 130 34 34	327 63 187 43 34	1,040 677 24 165 174	1,116 703 13 167 233	3,484 1,027 861 652 944	4,261 1,031 1,046 1,009 1,175	3,212 1,043 1,091 657 421	4,535 1,124 1,125 880 1,406
N. CENTRAL Dhio nd. II. Mich. Wis.	91 20 16 1 36	123 17 5 59 29 13	120 42 3 24 45 6	146 48 22 65 11	4,018 1,059 451 1,099 698 711	4,540 1,250 540 1,325 762 663	3,627 1,061 399 1,049 689 429	3,065 1,239 534 126 825 341
W.N. CENTRAL Minn. owa Mo. N. Dak. S. Dak. Nebr. Kans.	30 6 6 11	61 27 2 15 2 1 8	291 42 73 38 33 25 4 76	483 77 69 49 106 86 2 94	1,854 487 298 529 52 139 125 224	2,049 465 313 614 48 84 195 330	2,079 609 277 811 76 111	2,222 597 302 756 70 93 133 271
S. ATLANTIC Del. Md. D. C. Va. Va. W. Va. N. C. S. C. Ga. Fla.	239 101 13 44 1 16 6 12 44	277 5 94 15 47 4 32 2 19 59	1,902 30 279 399 123 508 103 294 166	2,032 47 355 486 103 493 142 268 138	7,337 79 694 72 1,152 113 1,134 768 1,258 2,067	6,705 105 684 56 849 139 940 641 1,204 2,088	4,977 87 770 U 747 124 954 627 1,210 458	5,126 115 603 U 811 133 983 493 1,509
E.S. CENTRAL Ky. Tenn. Ala. Miss.	31 12 11 6 2	42 17 11 13 1	184 26 96 60 2	181 19 92 69	2,238 328 546 617 747	2,044 329 538 570 607	1,618 210 663 474 271	1,560 225 699 522 114
W.S. CENTRAL Ark. La. Okla. Tex.	11 3 4 3	67 3 11 8 46	876 20 57 799	776 20 3 51 701	3,213 765 313 403 1,732	4,213 618 744 335 2,516	2,068 92 566 292 1,118	2,570 503 621 261 1,185
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	48 3 3 19 3 9 9	42 3 3 20 7 5 6	217 31 28 20 14 109 14	245 61 9 50 19 88 10 8	1,833 62 121 52 519 250 513 188 128	2,339 79 103 58 621 201 612 426 239	1,511 4 43 531 205 522 183 23	2,182 96 49 601 184 648 424 180
PACIFIC Wash. Oreg, Calif, Alaska Hawaii	185 9 11 155 1	233 25 36 163	309 3 269 37	282 7 249 26	4,086 442 207 3,084 34 319	4,673 487 259 3,669 53 205	3,408 491 271 2,335 28 283	4,429 577 315 3,297 33 207
Guam P.R. V.I. Amer, Samoa	3 U	2 5	80 Ú	67 Ú	459 U	23 572 U	מממט	U

N: Not notifiable. U: Unavailable. -: No reported cases.

* Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

* Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory information System (PHLIS).

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending October 27, 2001, and October 28, 2000 (43rd Week)*

-	NETS	Shigel		HLIS		ohilis Secondary)	Tube	rculosis
	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
Reporting Area INITED STATES	14,667	18,607	2001 6,786	10,688	2001 4,754	5.041	9,883	11,716
IEW ENGLAND Maine I.H. It. Mass. I.I. Conn.	228 6 6 7 177 177 15	355 10 6 4 249 26 60	245 2 3 5 170 23 42	339 11 8 226 30 64	49 1 2 27 9 10	74 1 2 52 4 15	340 8 14 4 198 29 87	353 16 17 4 204 27 85
MID. ATLANTIC Ipstate N.Y. I.Y. City I.J.	1,086 427 293 185 181	2,215 631 865 470 249	669 101 319 184 65	1,450 202 593 403 252	428 23 228 116 61	234 9 97 59 69	1,810 285 869 418 238	1,858 247 1,002 444 165
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	3,614 2,492 186 419 265 252	3,609 325 1,353 1,051 591 289	1,585 1,047 34 288 192 24	1,080 270 143 74 542 51	792 70 143 239 318 22	1,013 64 299 354 254 42	1,070 217 85 500 203 65	1,166 239 114 544 195 74
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	1,530 360 336 279 20 410 63 62	2,079 680 450 599 16 7 126 201	1,114 384 276 185 28 206	1,763 768 307 420 49 4 102 113	75 27 4 21 - 5 18	58 15 10 26 - 2 5	377 190 34 109 3 12 29	427 132 233 157 2 16 20 67
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga.	2,027 14 131 50 303 8 305 227 252 737	2,530 22 177 67 394 4 316 112 223 1,215	651 10 82 U 124 8 149 114 130 34	1,014 20 98 U 319 3 238 82 159 95	1,656 9 203 32 90 4 380 203 305 430	1,676 8 256 34 114 3 410 191 320 340	1,955 16 180 51 211 26 287 153 367 665	2,380 14 204 26 215 26 291 229 512 863
E.S. CENTRAL Ky. Tenn. Ala. Miss.	1,385 646 86 185 468	959 396 314 69 180	508 264 85 130 29	497 94 347 50 6	526 40 269 104 113	747 70 447 107 123	673 102 241 221 109	777 99 298 255 125
W.S. CENTRAL Ark, La. Okla, Tex.	1,944 497 121 66 1,260	2,912 174 241 107 2,390	1,098 155 137 17 789	926 51 151 38 686	604 31 138 57 378	697 91 185 101 320	758 127 119 512	1,710 158 146 128 1,278
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	806 5 34 3 210 110 331 48 66	1,019 7 44 5 221 134 423 72 113	1 239 72 236 47 8	741 25 3 180 98 293 76 66	203 1 1 36 18 131 8	199 1 1 8 15 168 1 5	402 6 8 3 99 24 178 32 52	428 14 7 2 71 36 174 41 83
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	2,047 174 76 1,734 6 57	2,929 401 152 2,337 7 32	313 167 91 6 49	2,878 371 98 2,377 3 29	421 42 13 356	343 55 11 276	2,498 199 84 2,065 40 120	2,617 211 83 2,121 88 114
Guam P.R. V.I. Amer. Samoa C.N.M.I.	8 U 4	35 29 U	ממממ	00000	234 U 4	130 U	76 U 23	119 U

N: Not notifiable. U: Unavailable. -: No reported cases.

Incidence data for reporting year 2001 are provisional and cumulative (year-to-date), Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending October 27, 2001, and October 28, 2000 (43rd Week)*

		ienzae,	1	lepatitis (V	iral), By Ty	pe			Measl	les (Rubeo	la)	
		sive	A		B		Indige	nous	Impo	rted'	Total	
Reporting Area	Cum. 2001 ^s	Cum. 2000	Cum. 2001	Cum. 2000	Cum. 2001	Cum. 2000	2001	Cum. 2001	2001	Cum. 2001	Cum. 2001	Cum. 2000
INITED STATES	1,046	1,065	8,391	10,977	5,286	5,827	-	49	120011	42	91	72
EW ENGLAND	78	84	540	330	83	94		4		1	5	6
Maine	2	1	10	19	5	5			+	- 2		
I.H.	4 3	12	16 14	18	14	15	-	1		-	1	3
Aass.	35	37	235	121	4	13		2	-	1	3	3
1.1.	5	4	55	22	25	18		-				- 2
onn.	29	23	210	141	31	37		1		-	1	-
MID. ATLANTIC	154	193	780	1,284	842	975		4	+	11	15	21
Ipstate N.Y.	57 38	82 52	210	210 443	110 346	109 476		1	-	4	5	10
N.Y. City	38	35	159	244	169	151	19	2		1	1	10
a.	20	24	180	387	217	239		1		5	6	1
N. CENTRAL	136	155	974	1,411	727	599				10	10	7
Ohio	49	47	171	227	84	93		-	+	3	3	2
nd.	43	26	92	89	42	41		*	-	4	4	
II. Mich.	10 12	53	373 282	614 408	124 477	104 326		-		3	3	3 2
Nis.	22	20	56	73	-	35	- 2	-	-	-		-
V.N. CENTRAL	54	63	351	595	169	249		4			4	1
Minn.	32	34	34	165	20	34	4	2	-		2	1
owa	13	19	31 97	61 241	21 90	30 122		2			2	-
Mo. N. Dak.	7	2	3	3	1	2		2			2	
S. Dak.	-	1	2	1	1	1					-	
Vebr.	1	3	30	30	19	37	U		U		9.	-
Kans.	1	4	154	94	17	23						
S. ATLANTIC Del.	308	237	1,963	1,215	1,163	1,034		4		1	5	4
Md.	74	73	230	175	121	110		2	-	7	3	
D.C.	-		43	23	11	28		-	-		:	-
Va. W. Va.	25 14	35 8	110	130 53	145	137		1			1	2
N.C.	44	21	193	123	173	208					~	
S.C.	6	7	66	70	28	14			- 2		3	
Ga. Fla.	72 73	57 36	752 552	243 384	305 360	181 330		1	- 3	7	1	2
E.S. CENTRAL	64	41	330	352	363	386		2			2	
Ky.	2	12	117	46	40	64		2	- 0	-	2	
Tenn.	34	17	129	123	195	180		8			0	-
Ala. Miss.	26	10	68 16	46 137	74 54	50 92		-	-	-		
	2	2					-		-			
W.S. CENTRAL Ark.	38	61	1,143	2,053 121	556 84	964 87		1	-	7	1	
La.	3	16	56	76	39	135						
Okla.	35	41	106	224	70	134			-		4	
Tex.		2	919	1,632	363	608	-	1	-		1	
MOUNTAIN	122	102	638	768	427	447	-	1		1	2	12
Mont. Idaho	1	4	10 53	23	11	6	-		-	1	1	
Wyo.		1	7	4	2	3						
Colo.	31	26	78	174	93	83	-	-	-	-	12.	- 2
N. Mex. Ariz.	20 54	20 35	33 342	65 386	124 128	121 167	- 1	1			1	
Utah	6	11	63	49	26	20			-	-		
Nev.	10	4	52	60	40	41	+	-		-		
PACIFIC	92	129	1,672	2,969	956	1,079		29	-	18	47	2
Wash.	4	5	125	245	117	93 97	-	13	-	2	15	3
Oreg. Calif.	17 43	29 33	1,462	154 2.544	91 723	868		10	-	11	21	14
Alaska	6	40	14	13	9	10	-			-		
Hawaii	22	22	3	13	16	11	-	2		5	7	3
Guam	- 5	1	-	1		9	U		U			
P.R. V.L	1	4	96	218	139	245	Ü	+	Ü	*		
Amer. Samoa	U	U	U	U	Ü	U	U	U	Ü	Ü	Ú	(
C.N.M.I.		Ü	-	Ü	28	Ü	U		U		-	i

C.N.M.I.

N: Not notifiable. U: Unavailable. -: No reported cases.

Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

For imported measles, cases include only those resulting from importation from other countries.

Of 223 cases among children aged <5 years, serotype was reported for 116, and of those, 19 were type b.

TABLE III. (Cont'd) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending October 27, 2001, and October 28, 2000 (43rd Week)*

		and	d Octo	ber 28,	2000	(43rd)	Week)*				
	Mening Dis	gococcal ease		Mumps			Pertussis			Rubella	
Reporting Area	Cum. 2001	Cum. 2000	2001	Cum. 2001	Cum. 2000	2001	Cum. 2001	Cum. 2000	2001	Cum. 2001	Cum. 2000
INITED STATES	1,748	1,841	3	182	276	35	3,794	5,679		20	146
NEW ENGLAND Maine N.H. At. Mass. R.I. Conn.	98 4 13 5 50 4 22	113 8 11 3 65 9			1 1 2	1	334 21 27 27 237 237 5	1,451 41 102 209 1,041 16 42			12 2 8 1
MID. ATLANTIC Upstate N.Y. V.Y. City V.J.	183 52 34 43 54	210 64 38 42 66		19 3 9 3 4	23 10 6 3 4	4	251 125 38 18 70	572 287 75 30 180		5 1 3 1	9 1 8
E.N. CENTRAL Dhia nd. III. Mich. Wis.	224 67 36 22 57 42	331 78 36 76 102 39		17 1 2 11 3	20 7 1 6 5	8 4 3 1	535 203 78 65 122 67	661 288 86 97 80 110		1 2	1
W.N. CENTRAL Minn. owa Mo. N. Dak. S. Dak. Nebr. Kans.	125 18 26 44 6 5 12	131 20 30 61 2 5 6	ŭ	7 3	17 7 4 1	2	245 105 20 89 4 4 4	477 284 46 74 6 7 24 36	Ü	3	1
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	326 4 37 35 12 60 31 40	256 1 26 37 12 34 21 43 82		34 6 - 6 - 5 5 7 5	40 9 9 6 10 2 4		31 1 36 2 63 31 14 26	430 8 110 3 97 1 94 26 36 55		6	94
E.S. CENTRAL Ky. Tenn. Ala. Miss.	119 20 56 30 13	122 25 51 33 13	2 2	8 3 1	5 1 2 2	1	128 34 55 36 4	103 52 31 17 3			6 1 1 4
W.S. CENTRAL Ark. La. Okla. Tex.	195 18 59 27 91	194 12 42 26 114		10 1 2 7	29 1 5	8 5 .	394 33 2 17 342	317 34 19 21 243		1	8 1 1 6
MOUNTAIN Mont. Idaho Wyo, Colo, N. Mex. Ariz, Utah Nev.	83 4 7 5 29 12 13 7 6	77 4 7 27 7 22 7 3		11 1 1 1 1 2 1 1 3	18 1 1 1 4 5 6	5	1.146 31 169 1 232 129 498 71 15	656 35 57 4 384 83 63 18		1	1
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	395 59 38 284 2	407 49 55 287 8 8	1 N 1	76 1 N 38 1 36	120 9 N 83 8	5 4 1	557 136 45 338 7 31	1,012 340 104 511 20 37	*****	1	13 7 6
Guam P.R. V.I. Amer. Samoa C.N.M.I.	4 U	9	ט י ט	Ü	14	ט טטט י	Ú	3 7 U	ט י ט	Ü	1 U

N: Not notifiable. U: Unavailable. -: No reported cases.

* Incidence data for reporting year 2001 are provisional and cumulative (year-to-date). Incidence data for reporting year 2000 are finalized and cumulative (year-to-date).

TABLE IV. Deaths in 122 U.S. cities,* week ending October 27, 2001 (43rd Week)

	1	All Cau	ses, By	Age (Y	ears)		P&I			All Cau	ses, By	Age (Y	(ears)		Parl
Reporting Area	All Ages	65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	-65	45-64	25-44	1-24	<1	Tota
NEW ENGLAND Boston, Mass. Gridgeport, Conn. Cambridge, Mass. Hartford, Conn. Lowell, Mass. Lynn, Mass. New Bedford, Mass. New Haven, Conn. Trovidence, R.I. Somerville, Mass. Springfield, Mass.	544 160 36 22 22 43 21 15 24 31 61 6	396 105 30 18 17 27 18 12 18 22 50 3	8	41 20 2 1 1 3 - 1 4 3 1 3	10 3 1 3 1 1	3 2	38 9 - 2 - 2 3 1 5 2	S, ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla. Miami, Fla. Norfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, Fl Tampa, Fla. Washington, D.C. Wilmington, Dec.	157	742 83 154 46 97 48 29 35 28 42 116 45 20	284 41 65 18 38 14 8 20 10 8 25 37	119 17 30 14 7 6 7 8 2 4 13	36 6 5 4 6 4 5 2	36 2 5 8 3 1 4 2 2 3 2 4	22
Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Allbany, N.Y. Allentown, Pa. Buffalo, N.Y. Camden, N.J. Eirzabeth, N.J. Erie, Pa.§	16 60 2,445 55 24 88 26 17 37	11,383 40 22,56 17 12,28	550 9 2 22 3 4 5	2 434 4 8 3 1	53 1 2 1 2 2	22 1	1 10 108 4 2 9 1	E.S. CENTRAL Birmingham, Ala, Chattanooga, Ten Konoxille, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Ala Nashville, Tenn.	734 151 69 94 U 163 96	491 94 45 64 U 110 65 28 85	163 41 19 19 U 34 19 8 23	46 10 2 7 U 9 7 2 8	17 4 3 U 4 3 1 2	17 1 3 1 U 6 2	3
Jersey City, N.J. New York City, N.Y. Newark, N.J. Philadelphia, Pa. Pittsburgh, Pa.S. Reading, Pa. Rochester, N.Y. Schenectady, N.Y. Scranton, Pa.S. Yyracuse, N.Y. Trenton, N.J. Juica, N.Y. Yonkers, N.Y.	51 1,493 U 33 247 36 15 145 27 32 68 23 28 U	42 679 U 19 167 27 13 121 20 25 52 17 26 U	U 11 59 5 1 18 5 5 9 3 1 1	386 U 10 2 1 6 2 1 5	1 32 U 1 8 1 1	15 U 1	40 U 2 8 1 16 5 2 9 3 3 3 U	W.S. CENTRAL Austin, Tex. Baton Rouge, La. Corpus Christi, To Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La. San Antonio, Tex Shreveport, La. Tulsa, Okla.	208 85 116 343 62 U	909 58 6 37 131 54 71 211 44 U 159 40 97	297 17 2 16 40 18 28 72 14 U 57 11 22	118 6 1 2 19 10 10 38 1 U 18 3	56 11 3 4 3 5 11 2 U 14	40 1 1 2 14 2 11 1 U 5 2	2 2 1 1
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, III. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind.	1,826 56 46 U 148 138 238 126 197 57	1,220 40 36 U 93 91 152 80 106 46	13 8 1 U 31 32 55 57 11	118 2 U 11 9 19 12 19	46 U 4 2 5 1 10	48 1 2 U 9 4 7 1 5 1	149 7 4 U 14 6 20 13 10 4 9	MOUNTAIN Albuquerque, N.I Boise, Idaho Colo. Springs, Co Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, Ut Tucson, Ariz.	36 101 101 191 20 159 36	687 81 28 39 56 129 15 95 29 89 126	183 23 5 8 18 42 4 31 6 21 25	71 4 1 4 11 11 11 21	44 6 2 4 11 6 4 5 6	20 1 5 3 8 1 2	1
Gary, Ind. Grand Rapids, Micl Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio	199 35 140 42 58 52 97	30 136 28 103 28 36 36 72 42	3 36 5 25 25 9 1 13 11 20 20	12 1 6 4 5 3	2 2 9 1 1 1 1 2 2 2	1 6 5 1 1 2	1 6 11 5 10 3 8 4 13	PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawai Long Beach, Calif. Los Angeles, Calif. Pasadena, Calif. Portland, Oreg. Sacramento, Calif.	f. 65 f. 322 33 111	1,168 11 77 12 59 44 212 24 74 100	345 4 19 5 16 16 68 5 22 38	103 4 1 6 4 23 3 3 6	36 2 2 1 11 11 6 4	35 1 1 8 6 3	13
W.N. CENTRAL Des Moines, lowa Duluth, Minn. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn Omaha, Nebr. St. Louis, Mo.	874 124 40 53 104 35 156 73	619 96 37 34 64 20 119	15 15 3 1 16 16 1 25 1 8 26 7 12	46 7 2 5 1 7 7	28 4 1 8 2 3 2 4	22 2 1 2 5 6	46 12 4 2 4 10 7	San Diego, Calif. San Francisco, Calif. San Jose, Calif. Santa Cruz, Calif. Seattle, Wash. Spokane, Wash. Tacoma, Wash. TOTAL	175 slif. 106 161	121 75 104 17 86 62 88	33 21 38 4 26 6 24	14 9 14 1 9 1 5	1 2 5 1	2 4 243	76
St. Paul, Minn. Wichita, Kans.	85 112	68	3 12	3 6	1 3	1 3	2 5								

U: Unavailable.

* Mortality data

Unavailable. ∹No reported cases.

Mortality data in this table are reported voluntarily from 122 cities in the United States, most of which have populations of ≥100,000. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included. Pneumonia and influenza.

Pneumonia and influenza.
 Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to 6 weeks.
 Total includes unknown ages.

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